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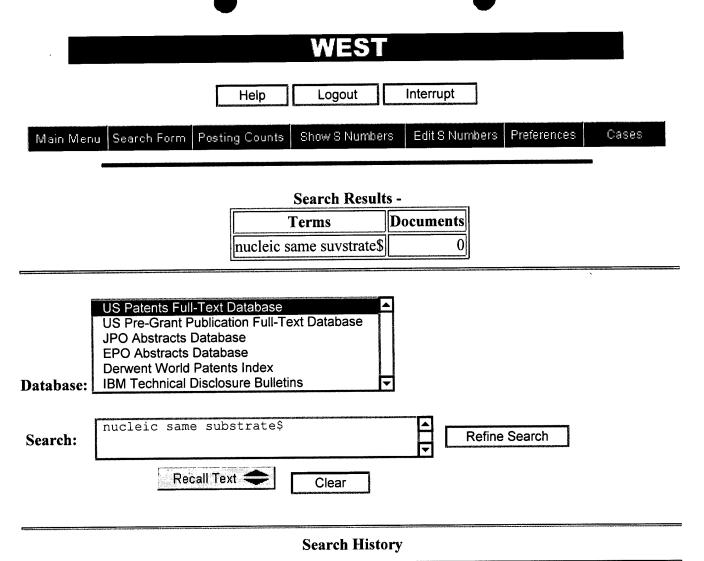
L12: Entry 1 of 3 File: USPT

DOCUMENT-IDENTIFIER: US 6284465 B1

TITLE: Apparatus, systems and method for locating nucleic acids bound to surfaces

Detailed Description Text (82):

The feasibility of locating features 13, 17 by a fluorescence signal 14 emitted from surface bound probes 12, which is different from the fluorescence signal 18 from a fluorophore labeled target sequence was demonstrated. Two different probe sequences 12 were synthesized in situ, using conventional techniques, but not directly labeled for the reason mentioned above for Example I experiment. Instead, two different target nucleotide sequences were biotinylated to indirectly label the probes 12 during hybridization. Streptavidin-labeled rhodamine-containing polystyrene microspheres were bound to the biotin, thereby providing the fluorescence label to the probes 12. The rhodamine absorbs green light and fluoresces in an orange-red color. The rhodamine microspheres were used in this experiment instead of the fluorescent dye pair microspheres mentioned above for Example 5. In either case, the feasibility of using fluorescence-containing microsphere labels was demonstrated. Also, two different target nucleotide sequences were labeled with a fluorophore Cy5 that absorbs red light and fluoresce in far red spectral region. The fluorescences from the Cy5 fluorophore label and the rhodamine are spectrally distinct for the purposes of the invention, in order to demonstrate measurement of both fluorescence channels (signals 14 and 18) from the same array.



DATE: Thursday, August 01, 2002 Printable Copy Create Case

Set Name side by side	Query	Hit Count	Set Name result set
DB=US	PT; PLUR=YES; OP=OR		
<u>L7</u>	nucleic same suvstrate\$	0	<u>L7</u>
<u>L6</u>	L5 same surface\$	3	<u>L6</u>
<u>L5</u>	11 same microsphere\$	7	<u>L5</u>
<u>L4</u>	L3 same (advantag\$ or useful\$)	2	<u>L4</u>
<u>L3</u>	L2 same hybridiz\$	30	<u>L3</u>
<u>L2</u>	L1 near0 nucleic	84	<u>L2</u>
<u>L1</u>	different near0 target\$	5375	<u>L1</u>

END OF SEARCH HISTORY

FILE 'HOME' ENTERED AT 12:07:40 ON 04 APR 2002)

	FILE	'MEDLIN	E, BIOSIS,	CAPLUS,	EMBASE'	ENTERED	ΑT	12:07:58	ON	04	APR	2002
L1		0 S	ARRAY SAM	E MICROS	PHERE?							
L2		196 S	ARRAY (P)	MICROSP:	HERE?							
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(FILE 'HOME' ENTERED AT 14:33:53 ON 01 AUG 2002)

	FILE 'MEDL	INE, BIOSIS, CAPLUS, EMBASE' ENTERED AT 14:34:09 ON 01 AUG 2002
L1		S FIBER(W)OPTIC(W)BUNDLE?
L2		S L1 (P) (NUCLEIC OR DNA OR POLYNUCLEOTIDE)
L3	0	S L2 (P) (ADVANTAG? OR USEFUL?)
L4	32	S L1 (P) (ADVANTAG? OR USEFUL?)
L5	1253325	S GLASS OR PLASTIC?
L6	9021	S L5 (P) (NUCLEIC OR DNA)
L7	1379	S L6 (P) HYBRIDIZ?
L8		S L7 (P) SUBSTRATE?
L9		S L8 (P) (ADVANTAG? OR USEFUL?)
L10	12	DUPLICATE REMOVE L9 (4 DUPLICATES REMOVED)
L11	0	S DECODER (W) LIGAND?
L12	5	S DECODER (P)LIGAND?
L13	2	S L12(P)(NUCLEIC OR DNA)

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English
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      The invention relates to compns. and methods for decoding microsphere
AB
      array sensors. It provides array compns. comprising a substate with a
      surface comprising discrete sites. The compn. further comprises a
      population of microspheres comprising at least a first and a second
      subpopulation; each subpopulation comprises a bioactive agent; and an
      identifier binding ligand that will bind a decoder
      binding ligand such that the identity of the bioactive agent can
      be elucidated. The microspheres are distributed on the surface. The
      microspheres comprise at least a first and a second subpopulation each
      comprising a bioactive agent and do not comprise an optical signature.
      The microspheres comprise at least a first and a second subpopulation
each
      comprising a bioactive agent and an identifier binding ligand
      that will bind a decoder binding ligand such that the
      identification of the bioactive agent can be elucidated. The invention
      provides methods of decoding an array compn. comprising providing an
      compn., and adding a plurality of decoding binding ligands to
      the array compn. to identify the location of at least a plurality of the
      bioactive agents. Bioactive agents are proteins or nucleic
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acids.